Claims

- 1.-19. (canceled)
- 20. (currently amended) A thermoplastic polymer blend comprising a thermoplastic starch, at least one synthetic polymer, and a hydrolysis component hydrolyzed polyvinyl acetate (PVAc) saponified to a hydrolysis degree of 20 to 70 %, and an acidic catalyst on PVAc basis, wherein the starch component of the polymer blend has a molecular weight which is only minimally reduced relative to native starch, wherein the thermoplastic polymer blend has a bi-continuous phase structure.
- 21. (original) The thermoplastic polymer blend according to claim 20, comprising extending agents, filling agents, internal lubricants, flow-improving agents, dyes, pigments, or mixtures thereof.
- 22. (currently amended) The thermoplastic polymer blend according to claim 20, comprising, relative to the total composition of the polymer blend, 30-70 % by weight of the thermoplastic starch, 20-40 % by weight of the synthetic polymer, and 6-25 % by weight of the hydrolysis hydrolyzed PVAc component on PVAc basis.
 - 23. (canceled)
- 24. (original) The thermoplastic polymer blend according to claim 20, wherein the synthetic polymer is a biologically degradable aliphatic polyester or a polyester copolymer or polyvinyl acetate or a polyvinyl acetate copolymer or a water-resistant starch derivative or a water-resistant cellulose derivative or polyvinyl alcohol or a polyvinyl alcohol copolymer.
- 25. (currently amended) A method for producing a thermoplastic polymer blend by reactive extrusion, the method comprising the steps of:
- a) mixing native starch, at least one hydrophobic polymer, a hydrolyzed component on polyvinyl acetate (PVAc) saponified to a hydrolysis degree of 20 to 70 % basis, and at least one of lower polyfunctional alcohols and water; and
- b) adding an acidic catalyst to the mixture of step a) and extruding the mixture in the presence of the acidic catalyst.
- 26. (original) The method according to claim 25, wherein the acidic catalyst is an organometallic compound selected from the group consisting of dibutyl tin oxide,

dibutyl tin dilaurate, tetra-2-ethylhexyl titanate, triethanolamine zirkonate, titanate compound chelated with lactic acid, triethanolamine titanate, and alkyl titanate.

- 27. (original) The method according to claim 26, wherein the mixture comprises 0.5 % to 2 % of the acidic catalyst, relative to the total weight of the mixture.
- 28. (original) The method according to claim 25, wherein the acidic catalyst is a Lewis acid.
- 29. (original) The method according to claim 28, wherein the mixture comprises 0.5 % to 2 % of the acidic catalyst, relative to the total weight of the mixture.
- 30. (original) The method according to claim 25, wherein the acidic catalyst is an acid selected from the group consisting of nitric acid, sulfuric acid, hydrochloric acid, and p-toluene sulfonic acid.
- 31. (currently amended) The method according to claim 30, wherein the mixture comprises 0.05 to 0.2 % of the acidic catalyst, relative to the its total weight of the mixture.
 - 32. (canceled)
- 33. currently amended) The method according to claim <u>25</u> [32], wherein the polyvinyl acetate is saponified to a hydrolysis degree of 30 % to 55 %.
- 34. (currently amended) The method according to claim <u>25</u> [32], wherein the polyvinyl acetate is prepared as an aqueous dispersion and is saponified at 120 140 °C with sodium hydroxide.
- 35. (currently amended) The method according to claim <u>25</u> [32], further comprising the step of adjusting the hydrolyzed component on polyvinyl acetate basis to a residual moisture contents of 15 35 %.
- 36. (currently amended) The method according to claim 25, wherein in the step a) the native starch, the hydrolyzed component on polyvinyl acetate basis, and the catalyst are mixed to a well-flowing powder mixture.
- 37. (original) The method according to claim 36, wherein approximately 1 % stearic acid, relative to the total weight of the powder mixture, is added to the powder mixture.
 - 38. (original) The method according to claim 36, wherein approximately 1 %

silica gel, relative to the total weight of the powder mixture, is added to the powder mixture.

39. (currently amended) The method according to claim <u>25</u> [32], wherein the acidic catalyst is metered in a liquid state mixed with glycerin.